Amendments to the Claims

This listing of claims will replace all prior versions and listing of claims in the application:

Listing of Claims

1. (Previously Presented) A compound having the following structure:

wherein

Z and Z_1 , are the same or different and are alkyl, aralkyl, aryl, aminoalkyl, alkyloxy, aralkyloxy, alkylamino, aralkylamino, arylamino, alkylmercaptan, aralkylmercaptan, arylmercaptan, carbohydrate, nucleoside, steroid, or substituted glyceride; and

X is methylene (-CH₂-), mono- or di-halo methylene.

- 2. (Original) The compound of claim 1, wherein Z and Z_1 are carbohydrates and X is methylene or difluoromethylene.
- 3. (Original) The compound of claim 1, wherein Z and Z_1 are nucleosides and X is methylene or difluoromethylene.
- 4. (Previously Presented) A method for the preparation of a compound having the following structure:

wherein

Z and Z_1 are the same or different and are alkyl, aralkyl, aryl, aminoalkyl, alkyloxy, aralkyloxy, alkylamino, aralkylamino, arylamino, alkylmercaptan, aralkylmercaptan, arylmercaptan, carbohydrate, nucleoside, steroid, or substituted glyceride; and

X is methylene (-CH₂-), mono- or di-halo methylene; which method comprises reacting a compound having the following structure:

wherein Z and X are as described, with a dehydrating agent.

- 5. (Original) The method of claim 4, wherein the dehydrating agent is a carbodiimide.
- 6. (Original) The method of claim 5, wherein the dehydrating agent is dicyclohexylcarbodiimide or diisopropylcarbodiimide.
- 7. (Original) The method of claim 4, wherein the starting material which is reacted with the dehydrating agent is selected from the group consisting of:
 - 2', 3'-O-isopropylideneadenosin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylideneinosin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylideneguanosin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylideneuridin-5'-ylphosphonomethylenephosphonate,
 - $2',\,3'\text{-}O\text{-}is opropylide necytidine-}5'\text{-}ylphosphonomethyle nephosphonate},$
 - 3'-O-(tetrahydropyranyl) thymidin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylidenetiazofurin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylidene-3-ribofuranosylbenzamid-5'-ylphosphono-methylenephosphonate,
 - 2', 3'-O-isopropylidene-Ψ-uridin-5'-ylphosphonomethylenephosphonate,
 - $\hbox{2', 3'-O-isopropylidene-Ψ-isocytidin-5'-ylphosphonomethylenephosphonate},$
- 9-(2'-deoxy-2'-fluoro-3'-O-tetrahydropyranyl-\(\beta\)-D-arabinofuranosyl) adenine-5-ylphosphonomethylenephosphonate,
- 9-(3'-deoxy-3'-fluoro-2'-O-tetrahydropyranyl ß-D-xylofuranosyl) adenine-5-ylphosphonomethylenephosphonate,

- 2'-deoxy-2'-fluoro-3'-O-tetrahydropyranyladenosin-5-ylphosphonomethylenephosphonate,
- $\label{lem:condition} 3'-deoxy-3'-fluoro-2'-O-tetrahydropyranyladenosin-5-ylphosphonomethylenephosphonate,$
 - 2', 3'-O-isopropylidene-9-deazaadenosin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylidene-9-deazainosin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylidene-9-deazaguanosin-5'-ylphosphonomethylenephosphonate,
 - 2', 3'-O-isopropylideneadenosin-5'-ylphosphonodifluoromethylenephosphonate,
 - 2', 3'-O-isopropylideneinosin-5'-ylphosphonodifluoromethylenephosphonate,
 - 2', 3'-O-isopropylideneguanosin-5'-ylphosphonodifluoromethylenephosphonate,
 - 3'-O-(tetrahydropyranyl) thymidin- 5'-ylphosphonodifluoromethylene-phosphonate,
 - 2', 3'-O-isopropylidenetiazofurin-5'-ylphosphonodifluoromethylenephosphonate,
- 2', 3'-O-isopropylidene-3-ribosylbenzamid-5'-ylphosphonodifluoromethylene-phosphonate,
 - 2', 3'-O-isopropylidene-Ψ-uridin-5'-ylphosphonodifluoromethylenephosphonate,
 - 2', 3'-O-isopropylidene-Ψ-isocytidin-5'-ylphosphonodifluoromethylene-phosphonate,
- $9\hbox{-}(2'\hbox{-}deoxy\hbox{-}2'\hbox{-}fluoro\hbox{-}3'\hbox{-}O\hbox{-}tetrahydropyranyl-}\beta\hbox{-}D\hbox{-}arabino fur an osyladen in e-5-ylphosphonodi fluoromethylene phosphonate,}$
- 9-(3'-deoxy-3'-fluoro-2'-O-tetrahydropyranyl-\(\beta\)-D-xylofuranosyl) adenine-5-ylphosphonodifluoromethylenephosphonate,
- $\label{lem:condition} 2'-deoxy-2'-fluoro-3'-O-tetrahydropyranyl-adenosin-5-ylphosphonodifluoro-methylenephosphonate,$
- 3'-deoxy-3'-fluoro-2'-O-tetrahydropyranyl-adenosin-5-ylphosphonodifluoro-methylenephosphonate,
 - 2', 3'-O-isopropylidene-9-deazaadenosin-5'-ylphosphonodifluoromethylene-phosphonate,
- $\label{eq:continuous} \hbox{2', 3'-O-isopropylidene-9-deazainosin-5'-ylphosphonodifluoromethylene-phosphonate,} \\$ and
- 2', 3'-O-isopropylidene-9-deazaguanosin-5'-ylphosphonoylphosphonodifluoro-methylenephosphonate.
- 8. (Previously Presented) A method for the preparation of a compound having the following structure:

wherein

Z and Z_1 are the same or different and are alkyl, aralkyl, aryl, aminoalkyl, alkyloxy, aralkyloxy, alkylamino, aralkylamino, arylamino, alkylmercaptan, aralkylmercaptan, arylmercaptan, carbohydrate, nucleoside, steroid, or substituted glyceride; and

X is methylene (-CH₂-), mono- or di-halo methylene; which method comprises reacting a compound having the following structure:

wherein Z, Z₁ and X are as defined above, with a dehydrating agent.

9. (Original) The method of claim 8, wherein the starting material which is reacted with the dehydrating agent is selected from the group consisting of:

P¹, P⁴-di (adenosin-5'-yl) phosphonomethylenephosphonic P², P³-anhydride,

 P^1 , P^4 -di [9- (2'-deoxy-2'-fluoro- β -D-arabinofuranosyl) -adenine-5-yl] phosphonomethylenephosphonic P^2 , P^3 -anhydride,

 P^1 , P^4 -di [9- (3'-deoxy-3'-fluoro- β -D-xylofuranosyl) -adenine-5'-yl] phosphonomethylenephosphonic P^2 , P^3 -anhydride,

 $P^1,\,P^4\text{-di (2'-deoxy-2'-fluoroadenosin-5-yl) phosphonomethylenephosphonic} \\ P^2,\,P^3\text{-anhydride},$

 $P^1,\,P^4\text{-di }(3'\text{-deoxy-3'-fluoroadenosin-5-yl})\ phosphonomethylenephosphonic \\ P^2,\,P^3\text{-anhydride},$

P¹, P⁴-di (inosin-5'-yl) phosphonomethylenephosphonic P², P³-anhydride,

 P^1 , P^4 -di (guanosi- n-5'-yl) phosphonomethylenephosphonic P^2 , P^3 -anhydride,

P¹, P⁴-di (uridin-5'-yl) phosphonomethylenephosphonic P², P³-anhydride,

- P^1 , P^4 -di (N^4 -acetylcytidin-5'-yl) phosphonomethylenephosphonic P^2 , P^3 -anhydride,
 - P¹, P⁴-di (thymidin-5'-yl) phosphonomethylenephosphonic P², P³-anhydride,
 - P¹, P⁴-di (tiazifurin-5'-yl) phosphonomethylenephosphonic P², P³-anhydride,
- P^1 , P^4 -di (3-ribosyl- benzamid-5'-yl) phosphonomethylenephosphonic P^2 , P^3 -anhydride,
 - P^1 , P^4 -di (Ψ -uridin-5'-yl) phosphonomethylenephosphonic P^2 , P^3 -anhydride,
 - P^1 , P^4 -di (Ψ -isocytidin-5'-yl) phosphonomethylenephosphonic
- P², P³-anhydride,
- P¹, P⁴-di (9-deazaadenosin-5'-yl) phosphonomethylenephosphonic P², P³-anhydride,
- P^1 , P^4 -di (9-deazainosin-5'-yl) phosphonomethylenephosphonic P^2 , P^3 -anhydride,
- P^1 , P^4 -di (9-deazaguanosin-5'-yl) phosphonomethylenephosphonic P^2 , P^3 -anhydride,
- P¹, P⁴-di (adenosin-5'-yl) phosphonodifluoromethylenephosphonic P², P³-anhydride,
- $P^1,\,P^4\text{-di (inosin-5'-yl) phosphonodifluoromethylene phosphonic} \\ P^2,\,P^3\text{-anhydride},$
- P^1 , P^4 -di (guanosin-5-yl) phosphonodifluoromethylenephosphonic P^2 , P^3 -anhydride,
- P^1 , P^4 -di (thymidin-5'-yl) phosphonodifluoromethylenephosphonic P^2 , P^3 -anhydride,
- $P^1,\, P^4\text{-di (tiazofurin-5'-yl) phosphonodifluoromethylenephosphonic} \\ P^2,\, P^3\text{-anhydride},$
- $P^1,\,P^4\text{-di }(3\text{-ribosylbenzamid-5'-yl})\ phosphonodifluoromethylenephosphonic}$ $P^2,\,P^3\text{-anhydride},$
- $P^1,\,P^4\text{-di}\,(\Psi\text{-uridin-5'-yl})$ phosphonodifluoromethylenephosphonic $P^2,\,P^3\text{-anhydride},$
- P^1 , P^4 -di (Ψ -isocytidin-5'-yl) phosphonodifluoromethylenephosphonic P^2 , P^3 -anhydride,
 - P¹, P⁴-di (9-deazaadenosin-5'-yl) phosphonodifluoromethylenephosphonic

- P², P³-anhydride,
 - P¹, P⁴-di (9-deazainosin-5'-yl) phosphonodifluoromethylenephosphonic
- P², P³-anhydride, and
- P^1 , P^4 -di (9-deazaguanosin-5'-yl) phosphonodifluoromethylenephosphonic P^2 , P^3 -anhydride.
- 10. 26. (Cancelled)
- 27. (New) The method of claim 4, additionally comprising reacting the product compound of claim 4 with a nucleophilic agent.
- 28. (New) The method of claim 27, wherein the nucleophilic agent is a compound comprising an –OH, -SH, -NH₂, or –NH moiety.
- 29. (New) The method of claim 27, wherein the nucleophilic agent is aralkyl, aryl, aminoalkyl, alkyloxy, aralkyloxy, alkylamino, aralkylamino, arylamino, alkylmercaptan, aralkylmercaptan, or arylmercaptan, carbohydrate, nucleoside, a mycophenolic acid residue or derivative, steroid, or substituted glyceride;
- 30. (New) The method of claim 27, wherein the nucleophilic agent is water.
- 31. (New) The method of claim 8, additionally comprising reacting the product compound of claim 8 with a nucleophilic agent.
- 32. (New) The method of claim 31, wherein the nucleophilic agent is a compound comprising an –OH, -SH, -NH₂, or –NH moiety.
- 33. (New) The method of claim 31, wherein the nucleophilic agent is water, alkyl, aralkyl, aryl, aminoalkyl, alkyloxy, aralkyloxy, alkylamino, aralkylamino, arylamino, alkylmercaptan, aralkylmercaptan, or arylmercaptan, carbohydrate, nucleoside, a mycophenolic acid residue or derivative, steroid, or substituted glyceride;

34.	(New)	The method of claim 31, wherein the nucleophilic agent is water.